

## Science Applications International Corporation

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April 8, 1991

REF: RFEV24-SAG-EGRF-T-053

TO:

Olga Erlich

LOCATION: EG&G RF

FROM:

Larry McInroy/Phil Ralphs

LOCATION: SAIC/Golden

SUBJECT:

Response to the Environmental Protection Agency (EPA) and the

Colorado Department of Health (CDH) Review Comments on the Draft

Site-Wide Treatability Studies Plan

In response to your request, please find attached our responses to the EPA and CDH review comments on the subject document that pertain to quality assurance. A handwritten response to these comments was faxed to you earlier. If you have any questions regarding these responses, please contact us at 279-7242

APPROVED

**EBASCO** 

DATE.

cc

T Greengard M Brooks

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Providing Waste Management Services for EG&G Rocky Flats Inc.

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ROCKY FLATS ENVIRONMENTAL RESTORATION PROGRAM REVIEW/COMMENT RESOLUTION FORM	J Title	Name(s) Environmental Protection Agency and Colorado Department of Health Date 4/8/91	REVIEWER'S COMMENTS REVIEWER'S COMMENTS	PAGE COMMENTS DISCUSSION	Several sections of the TSP address the question of compliance with and preparation of other program documents in many instances the terminology used is inconsistent and the text provided indicates general confusion over how all these documents fit together. These passages must be revised in accordance with verbal comments provide for the TSP and written comments on the SOP/QAPJP framework, and test-specific documentation (SOPAs/QAAs) provided as needed	The goal of treatability studies, as will be stated in individual TSPs, is to defined in the TSP. The basis for evaluating the effectiveness of a treatment technology is to calculate removal efficiencies and compliance with established cleanup criteria. Similarly, data quality needs of the treatment established in the TSP to define the data quality needs of the treatability studies program studies program at the TSP to define the data quality needs of the treatability assurance Addenda that will be developed for each TSP. The relationship between the QAPIP and QAAs is explained in this section.	Treatability tests are required to ensure the data collected are accurate, complete and appropriate A usual objective in data collection is to obtain measurements and samples that meet acceptable standards of accuracy, precision, comparability, representativeness, and completeness. The taxt studies Also the data quality objectives for site-wide treatability studies about the data quality objectives for site-wide treatability studies.
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	Document	Reviewer Name(s)		COMMENT	<del>-</del>	7	m

ROCKY FLATS ENVIRONMENTAL RESTORATION PROGRAM REVIEW/COMMENT RESOLUTION FORM	Page 2 of 2	and Colorado Department of Health 4/8/91	RESPONSE	DISCUSSION	As is now stated in Section 6 2 11, a Quality Assurance Addende (QAA) to the Site-Wide Quality Assurance Project Plan (QAP)P) will be developed for each TSP. The Site-Wide SAP, which consists of the SOPs and QAP)P, will not be modified, since, as you correctly point out, these are generic documents that address all IAG activities.  I to do work  I work  I by  Site-Wide SAP, which consists of the SOPs and QAP)P, will not be modified, since, as you correctly point out, these are generic documents that address all IAG activities  By  Site Wide SAP, which compared to EPA and CDH for review and approval. These SOPs and the QAPP together comprise the SAP required by the IAG The SOPs include field sampling procedures and appropriate QAPQC procedures. Additional QA/QC controls are included in the QAPP. Where the implementation of a treatability study requires the use of an SOP or QA/QC control described in the QAPP, will be referenced, but not included, in either the work plan procedures that are specific to the treatability study will be included in the Work Plan.	
	and Title Draft Site-Wide Treatability Studies Plan	Environmental Protection Agency	REVIEWER'S COMMENTS	COMMENTS	Section 3.0, page 3.2. Any specific field or quality assurance activities required to conduct the treatability studies should be incorporated by the use of mechanism that does not require modification of the Site Wide Sampling and Analysis Plan (SAP) or the Site-Wide Quality Assurance Project Plan (QAPJP), since these are generic documents. One way to do this, is to include the required addenda within the treatability study work plans and then incorporate them into the Site-Wide SAP and QAPJP by reference.  The FSP and QAPJP are not conducted within the site-wide treatability studies program and they should not be modified to meet the needs of each treatability study report for use as a basic reference document in the dorumity Study report for use as a basic reference document in the completion of Feasibility Studies.  Section 6.1, page 6.1. FSP and QA/QC procedures specific to the treatability studies to be conducted must be included in the treatability studies work plan and as agenda to the generic FSP and QA/QC procedures.	
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